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FORM P (REV. 1	1-2000)	PARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY'S DOCKET NUMBER					
'	TRANSMITTAL LETTER T	TO THE UNITED STATES	0224 0421 D					
	DESIGNATED/ELECTE	D OFFICE (DO/EO/US)	0234-0421P U.S. APPLICATION NO, (If known, see 37 CFR 1.5)					
	CONCERNING A FILING		09 MEV8 6626					
INTE	RNATIONAL APPLICATION NO.	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED					
	DC# (TD00 (01050							
TETT I	PCT/JP00/01959 E OF INVENTION	March 29, 2000	December 13, 1999					
		CTION OF HIGH-FUNCTION PHOTOCAT	AT VOT					
APPL	ICANT(S) FOR DO/EO/US	or and remersion inclocal	ALISI					
*	TANAKA, Keiichi; VOHRA, Muhammad Sharing Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:							
Applic	ant herewith submits to the United States	Designated/Elected Office (DO/EO/US) the follo	owing items and other information:					
1.	This is a FIRST submission of items conce	erning a filing under 35 U.S.C. 371.						
2	This is a SECOND or SUBSEQUENT sub	omission of items concerning a filing under 35 U.S.	C. 371.					
3.	This express request to begin national	examination procedures (35 U.S.C. 371(f)) at a	any time rather than delay					
K 7	examination until the expiration of the	applicable time limit set in 35 U.S.C. 371(b) a	and PCT Articles 22 and 39 (1).					
4. 🔀	The US has been elected by the expiration of 19	months from the priority date (Article 31).						
5. 🔀	A copy of the International Application							
, m = 1,		d only if not transmitted by the International E	Bureau).					
F. S.	b. has been transmitted by the Inte	ernational Bureau.						
74 711 —		on was filed in the United States Receiving Of						
6. D	An English language translation of the	ne International Application as filed (35 U.S.C	. 371(c)(2)).					
M	a. is transmitted herewith.							
NJ_	b. has been previously submitted in	under 35 U.S.C. 154(d)(4)						
7. 📉	Amendments to the claims of the Inter	national Application under PCT Article 19 (33	5 U.S.C. 371(c)(3)).					
# #. z.	a. are transmitted herewith (requir	ed only if not transmitted by the International	Bureau).					
F17	b. have been transmitted by the In							
We will limit with	c. have not been made; however, t	he time limit for making such amendments ha	s NOT expired.					
	d. Mave not been made and will no		_					
8.	An English language translation of the	e amendments to the claims under PCT Article	e 19 (35 U.S.C. 371(c)(3)).					
9.	An oath or declaration of the inventor	(s) (35 U.S.C. 371(c)(4)).						
10.	An English language translation of the	e annexes of the International Preliminary Exa	mination Report under PCT Article 36					
ems 1	(35 U.S.C. 371(c)(5)).		-					
	1. to 20. below concern document(s)	or information included:						
11.	An Information Disclosure Statement	under 37 CFR 1.97 and 1.98-International Sea	arch Report (PCT/ISA/210) with documents					
12.		g. A separate cover sheet in compliance with 3	37 CFR 3.28 and 3.31 is included.					
13.	A FIRST preliminary amendment.							
14.	A SECOND or SUBSEQUENT prelin	ninary amendment.						
15.	A substitute specification.							
16.	A change of power of attorney and/or							
17.	A computer-readable form of the sequ	ence listing in accordance with PCT Rule 13te	er.2 and 35 U.S.C. 1.821-1.825.					
18.	A second copy of the published interna-	ational application under 35 U.S.C. 154(d)(4).						
19.	A second copy of the English language	e translation of the international application un	nder 35 U.S.C. 154(d)(4).					
20. 🔀	Other items or information:							
	 PCT Request (PCT/RO/101) Verification of Translation 							
	3.) Zero (0) sheets of Formal Drawings	S						

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21. The following fees are submitted:					CAL	CULATIONS	PTO USE ONLY
BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5):							
	Neither international preliminary examination fee (37 CFR 1.482)						
nor international search				\$1,000.00			
and international Sear	cn Report not prepare	d by the E	PO or JPO	\$1,000.00			
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Independent Claims	2 - 3 =		0	X \$80.00	\$	0	
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	NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.						
Send all correspondence to	o:						
Birch, Stewart, Kola	Birch, Stewart, Kolasch & Birch, LLP or Customer No. 2292						
P.O. Box 747						_	
Falls Church, VA 22040-0747							
(703)205-8000					1		
Date: March 7, 2001						, who	
				Marc	S. We	iner, #32,181	
/cqc							
							

VERIFICATION OF TRANSLATION

RE: INTERNATIONAL APPLICATION NO. PCT/JP00/01959

I, Toshizo Iida of ISHII Bldg. 3F, 1-10, Shimbashi 3-chome, Minato-ku, Tokyo 105-0004 Japan, am the translator of the documents attached and I state that the following is a true translation to the best of my knowledge and belief.

Signature of translator

Dated: March 6, 2001

Toshizo IIDA

09/786626

JC02 Rec'd PCT/PTO 0421P MAR 2001

IN THE U.S. PATENT AND TRADEMARK OFFICE

Applicant:

TANAKA, Keiichi et al.

Int'l. Appl. No.: PCT/JP00/01959

Appl. No.:

New

Group:

Filed:

March 7, 2001

Examiner:

For:

PRODUCTION OF HIGH-FUNCTION PHOTOCATALYST

PRELIMINARY AMENDMENT

BOX PATENT APPLICATION

Assistant Commissioner for Patents Washington, DC 20231

March 7, 2001

Sir:

following Preliminary Amendments and Remarks respectfully submitted in connection with the above-identified application.

AMENDMENTS

IN THE SPECIFICATION:

Please amend the specification as follows:

Before line 1, insert -- This application is the national phase under 35 U.S.C. § 371 of PCT International Application No. PCT/JP00/01959 which has an International filing date of March 29, 2000, which designated the United States of America. --

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IN THE CLAIMS:

Please amend the claims as follows:

- 4. (Amended) The high-function photocatalyst according to claim 1 or 2, wherein the polymer having an anionic group is poly(fluorine-substituted sulfonic acid).
- 5. (Amended) The high-function photocatalyst according to claim 1 or 2, wherein the photocatalyst is spherical.
- 6. (Amended) The high-function photocatalyst according to claim 1 or 2, wherein the photocatalyst is immobilized on a substrate.

MSW/cqc

0234-0421P

REMARKS

The specification has been amended to provide a crossreference to the previously filed International Application. claims have also been amended to correct improper multiple dependencies and to place the application into better form for examination. Entry of the present amendment and favorable action on the above-identified application are earnestly solicited.

If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies, to charge payment or credit any overpayment to Deposit Account No. 02-2448 for any additional fees required under 37 C.F.R. § 1.16 or under 37 C.F.R. § 1.17; particularly, extension of time fees.

Respectfully submitted,

BIRCH, STEWART, KOLASCH & BIRCH, LLP

Marc S. Weiner,

P.O. Box 747

Falls Church, VA 22040-0747

(703) 205-8000

Attachment: Version with Markings to Show Changes Made

(Rev. 01/22/01)

VERSION WITH MARKINGS TO SHOW CHANGES MADE

The specification has been amended to provide a cross-referencing paragraph to the International Application.

The claims have been amended as follows:

- 4. (Amended) The high-function photocatalyst according to [any one of claims 1 to 3]claim 1 or 2, wherein the polymer having an anionic group is poly(fluorine-substituted sulfonic acid) [(for example, Nafion)].
- 5. (Amended) The high-function photocatalyst according to [any one of claims 1 to 4] claim 1 or 2, wherein the photocatalyst is spherical.
- 6. (Amended) The high-function photocatalyst according to [any one of claims 1 to 5] claim 1 or 2, wherein the photocatalyst is immobilized on a substrate.

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SPECIFICATION

PRODUCTION OF HIGH-FUNCTION PHOTOCATALYST

5 TECHNICAL FIELD

The present invention relates to a photocatalyst used, for example, in decomposition of harmful organic compounds, a method of manufacturing the same, and a photocatalyst that is obtained immobilizing said photocatalyst.

BACKGROUND ART

Wastewater that can be processed by a photocatalyst at a practical level is limited. This is because the processing efficiency of a current photocatalyst is not sufficient with regard to most harmful substances. To improve the efficiency, it is considered to carry platinum on the photocatalyst or dope with impurities, but the effect is inadequate, and stable results are not obtained in the latter method.

It is hence an object of the present invention to provide a high-function photocatalyst that exhibits high decomposition efficiency toward harmful materials and that can be used for a long time, and a method of manufacturing

the same.

Other and further objects, features, and advantages of the invention will appear more fully from the following description.

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DISCLOSURE OF THE INVENTION

The present inventors, after intensive studies to solve the above problems, discovered that many harmful substances have a positive electric charge in water.

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To decompose such harmful substances having a positive electric charge efficiently, it has been found that the photocatalyst should be brought into as closely to the harmful substances as possible, and hence the present invention has been completed.

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That is, the present invention was accomplished based on the findings that a significant photocatalytic function is expressed toward harmful materials having a positive electric charge, when the surface of a spherical photocatalyst is partially covered with a polymer having an anionic group.

20 an anionic group.

In the present invention, to cover the surface of a spherical photocatalyst partially with a polymer having an anionic group, the polymer having an anionic group is dissolved in a solvent, to dilute the concentration of the

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polymer, and then the solution is evaporated and dried, so that the polymer can be entangled in parts of the surface of the spherical photocatalyst.

If the entire photocatalyst is covered with the polymer having an anionic group, chances of light and water existing simultaneously around the catalyst are lowered, and hence it is required that the catalyst should be exposed partially.

10 BEST MODE FOR CARRYING OUT THE INVENTION

The photocatalyst utilized in the present invention may be any photocatalyst that can be ordinarily used, and it is not particularly limited. Specific examples of the photocatalyst include titanium dioxide, zinc oxide,

2 zirconium oxide, and tungsten oxide and the like, among them, titanium dioxide being preferred.

The photocatalyst may be used in a form of powder, immobilized powder or film prepared by sol-gel method or vapor deposition method. The shape of the photocatalyst is not particularly limited, and spherical, flat, tubular or fibrous shape may be used.

The photocatalyst, the polymer having an anionic group, and the solvent are mixed, stirred and dried, or alternatively the photocatalyst is immobilized on a glass plate, or natural or synthetic high polymer film or the

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like, and a solution dissolving the polymer having an anionic group is applied and dried thereon. For example, the photocatalyst may be immobilized with a binder on the ceramic or plastic film. A diluting solvent of the polymer having an anionic group is not particularly limited if it may be any solvent capable of dissolving this polymer, and for example, methanol, ethanol, propanol and the like may be used.

The polymer having an anionic group includes

10 poly(fluorine-subsutituted sulfonic acid) such as Nafion
(tradename of Du Pont Company), poly(fluorine containing
carboxylic acid) such as Flemion (tradename of Asahi Glass
Company), polystyrene sulfonic acid, polyvinyl sulfonic
acid and the like, and, among them, Nafion is particularly
15 preferred because of its strong resistance toward
decomposition of photocatalyst.

A linear polymer is preferred, and it is required to be insoluble in water and soluble in organic solvent, and the molecular weight is preferably about 500,000 to 1,000,000.

The amount to be used of the polymer having an anionic group is preferred to be 0.05 to 5 ml in a 5-% by weight solution per g of photocatalyst powder, and more preferably 0.1 to 0.4 ml. In the immobilized

25 photocatalyst, it is preferred to be 0.1 to 1 ml of 5-% by

weight solution per surface area of 20 cm², more preferably 0.1 to 0.3 ml. The solution is mixed and applied to be uniform, dried at room temperature. Thus, a partial covering can be formed.

5 In the present invention, the surface of the photocatalyst is partially covered with the polymer having an anionic group. Herein, the partial covering means covering of the surface of the photocatalyst with the polymer so that at least a part of the photocatalyst surface may be exposed so as not to impede the catalytic 10 function of the photocatalyst, and to attract the organic materials having positive ions around the catalyst enough electrostatically, by the anionic group of the polymer existing on the photocatalyst surface. In the present 15 invention, the amount for use of polymer having an anionic group, used for partially covering the surface of the photocatalyst differs according to the concentration of the organic materials to be decomposed or the type of the photocatalyst, but may be set properly within the specified range depending on the situation. 20

The photocatalyst of the present invention is effective for harmful materials having positive ions. For example, amine compound, imine, pyridine compound and their salts and the like are particularly effective.

Using the photocatalyst of the present invention,

these compounds contained in water can be decomposed at high efficiency. The decomposition process can be conducted by that the wastewater to be treated is brought into contact with the photocatalyst, and irradiated with ultraviolet ray.

Applicable objects of the photocatalyst of the present invention are not limited to harmful substances in water, but include harmful gases, for example, gaseous amine.

The light source of irradiation is, preferably, a light source containing light of shorter wavelength than 380 nm. Such examples include low pressure or high pressure mercury vapor lamp, xenon lamp, halogen lamp, blacklight, and sunlight etc.

15 The mechanism why the photocatalyst of the present invention can decompose the organic materials having a positive ion efficiently is not fully understood, but it seems that the anionic group of the polymer existing on the surface of the photocatalyst attracts the organic materials having a positive ion to the photocatalyst, and that the hydroxyl radical released by the photocatalyst at a close distance efficiently attacks the organic materials having a positive ion.

The embodiments of the present invention may be summarized as follows.

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- (1) A high-function photocatalyst having its surface partially covered with a polymer having an anionic group.
- (2) The high-function photocatalyst according to above (1), wherein the polymer is a linear polymer.
- (3) The high-function photocatalyst according to above (1) or (2), wherein the photocatalyst is in a form of a fine powder with particle size of 0.04 to 1 μm .
- (4) The high-function photocatalyst according to any one of above (1) to (3), wherein the polymer having an anionic group is poly(fluorine-substituted sulfonic acid) (for example, Nafion).
- (5) The high-function photocatalyst according to any one of above (1) to (4), wherein the photocatalyst is spherical.
- 15 (6) The high-function photocatalyst according to any one of above (1) to (5), wherein the photocatalyst is immobilized.
 - (7) A method of manufacturing a high-function photocatalyst comprising the steps of adding a spherical photocatalyst into a solution having a linear polymer having an anionic group dissolved in a solvent, stirring, and drying.
 - (8) A method of manufacturing a high-function photocatalyst comprising the steps of immobilizing a photocatalyst on a substrate of a film or the like, for

example, with an adhesive, applying thereon a solution dissolving a polymer having an anionic group, and drying.

The photocatalyst of the present invention has been confirmed to decompose organic materials having a positive electric charge efficiently. The efficiency of the photocatalyst itself is also confirmed not to deteriorate for a long period.

Examples

Next, the present invention will be described in more detail based on examples given below, but the present invention is not meant to be limited by the following examples.

Example 1

15 To 0.2 ml of 5-% by weight commercial Nafion solution, 1 ml of methanol was added, and 2 g of titanium dioxide powder (mean particle size 0.15 μm) was mixed to the solution, and the mixture was dried overnight at room temperature. In 500 ml of 10⁻⁴ mol 1⁻¹ (26 ppm) solution of herbicide paraquat, 2 g of titanium dioxide covered with the above-described Nafion was suspended. The suspension was irradiated with a blacklight of 6 W placed in the center of the liquid. First, by stirring the suspension for 120 minutes without irradiation with light, 10% of the initial concentration of the herbicide was

decreased. Then, starting irradiation with the light, 75% of the concentration was decomposed after 20 minutes, and 100% of the concentration was decomposed in 90 minutes. In a comparative example of titanium dioxide without covering with Nafion, only 25% of the concentration was decomposed in 20 minutes and 55% of the concentration in 60 minutes.

Example 2

The same experiment as in Example 1 was conducted

10 except that 2 ml of Nafion solution was used with 2 g of
titanium dioxide. First, by stirring for 120 minutes
without irradiation with light, 50% of the initial
concentration of the herbicide was decreased. It is
considered the result was due to adsorption. Later, by

15 irradiation with the light for 5 minutes, only 3% of the
concentration was detected.

Example 3

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The same experiment as in Example 1 was conducted except that ethylamine of 10⁻⁴ mol 1⁻¹ (6.9 ppm) was used instead of paraquat. By irradiation with the light for 5 minutes, 55% of the initial concentration of the ethylamine was decomposed, and 80% of the concentration was decomposed in 10 minutes. In case of the titanium dioxide without covering with Nafion, almost nothing of the concentration was decomposed in the first 5 minutes,

and only 20% of the concentration was decomposed after 10 minutes.

Example 4

0.2 ml of 5-% by weight commercial Nafion solution was diluted with 0.5 ml of methyl alcohol, and the solution was applied uniformly on a titanium dioxide thin film prepared by sol-gel method on a glass plate of 45 x 45 mm, and the film was dried for 24 hours at room temperature. This dried plate was put into a cell made of Pyrex glass of 50 (width) x 50 (length) x 10 (thick) mm, 10 and 15 ml of paraquat solution of 10^{-4} mol 1^{-1} was added therein, and the cell was irradiated with high pressure mercury vapor lamp of 500 W. First, by stirring for 90 minutes without irradiation with light, 10% of the initial concentration of the herbicide was decreased. Then, by 15 irradiation with the light, 75% of the initial concentration was decreased in 60 minutes. Example 5

To investigate the stability of covering Nafion film,

the same experiment as in Example 1 was conducted by using deionized water instead of paraquat. In 27 hours without irradiation with light, 4.5 x 10⁻⁵ mol 1⁻¹ of sulfuric acid ion and 3 ppm of TOC were detected, and there was almost no change of these concentrations until the end of 51 hours. It was considered that sulfuric acid ions were

derived from titanium dioxide, and TOC were derived from impurities in Nafion. Afterwards, for 19 days consecutively, irradiation with the light was continued, and samples were taken at proper time intervals. There was no change in sulfuric acid ions, but TOC decreased slightly. Within this time duration, the results suggest that Nafion is stable.

Example 6

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To evaluate reproducibility of the above-described photocatalyst when used repeatedly, after the experiment of example 1, the photocatalyst in the suspension was collected, and a new solution of paraquat was added therein, and the irradiation to the suspension was conducted in the same condition. This operation was repeated 5 times, and the decomposition efficiency was measured each time, but deterioration of efficiency of photocatalyst was not detected.

INDUSTRIAL APPLICABILITY

20 The photocatalyst of the present invention is preferable as photocatalyst for decomposing organic materials having positive electric charge efficiently.

Having described our invention as related to the present embodiments, it is our intention that the

invention not be limited by any of the details of the description, unless otherwise specified, but rather be construed broadly within its spirit and scope as set out in the accompanying claims.

CLAIMS

- 1. A high-function photocatalyst having its surface partially covered with a polymer having an anionic group.
- 5 2. The high-function photocatalyst according to claim 1, wherein the polymer is a linear polymer.
 - 3. The high-function photocatalyst according to claim 1 or 2, wherein the photocatalyst is in a form of a fine powder with particle size of 0.04 to 1 μm .
- 4. The high-function photocatalyst according to any one of claims 1 to 3, wherein the polymer having an anionic group is poly(fluorine-substituted sulfonic acid) (for example, Nafion).
- 5. The high-function photocatalyst according to any one of claims 1 to 4, wherein the photocatalyst is spherical.
 - 6. The high-function photocatalyst according to any one of claims 1 to 5, wherein the photocatalyst is immobilized on a substrate.
- 7. A method of manufacturing a high-function photocatalyst comprising the steps of adding a spherical photocatalyst into a solution having a linear polymer having an anionic group dissolved in a solvent, stirring, and drying.
- 8. A method of manufacturing a high-function

photocatalyst comprising the steps of immobilizing a photocatalyst on a substrate of a film or the like, applying thereon a solution dissolving a polymer having an anionic group, and drying.

ABSTRACT OF THE DISCLOSURE

A high-function photocatalyst having its spherical surface partially covered with a polymer having an anionic group is disclosed. This photocatalyst has high efficiency in decomposition of harmful substances, and it is a high-function photocatalyst usable for a long period of time.

BIRCH, STEWART, KOLASCH & BIRCH, LLP

COMBINED DECLARATION AND POWER OF ATTORNEY

PLEASE NOTE: YOU MUST COMPLETE THE FOLLOWING:

FOR PATENT AND DESIGN APPLICATIONS

ATTORNEY DOCKET NO. 234–421P

(Status - patented, pending, abandoned)

As a below named inventor, I hereby declare that: my residence, post office address and citizenship are a
stated next to my name; that I verily believe that I am the original, first and sole inventor (if only one invento
is named below) or an original, first and joint inventor (if plural inventors are named below) of the subject
matter which is claimed and for which a patent is sought on the invention entitled:

Insert Title:	is named below) or an original matter which is claimed and for PRODUCTION OF	, first and joint inventor r which a patent is sought HIGH-FUNCTION PI	t on the invention entitled:	ed below) of the subject
Fill in Appropriate Information -	the specification of which is att	ached hereto. If not atta was filed on		as
For Use Without Specification		ation Number		; and /or
Attached:	the enecification	was filed onMa	arch 29, 2000	as PCT
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	amended under PC	Γ Article 19 on	(if a	
The state of the s	including the claims, as amenda I acknowledge the duty to Code of Federal Regulations, § I do not know and do not I my or our invention thereof, or our invention thereof or more on sale in the United States of a been patented or made the subcountry foreign to the United assigns more than twelve monta patent or inventor's certificated.	led by any amendment re- disclose information whis \$1.56. believe the same was ever learn patented or described in than one year prior to the America more than one yea spect of an inventor's certicates of America on an a hs (six months for designs to on this invention has been on by me or my legal rep- tiority benefits under Title pyentor's certificate liste	known or used in the United any printed publication in any printed publication in a application, that the same ar prior to this application, ficate issued before the date application filed by me or mean prior to this application, a cen filed in any country foreign resentatives or assigns, exceeds 35, United States Code, § d below and have also ide	lity as defined in Title 37, d States of America before any country before my or the was not in public use or that the invention has not to of this application in any my legal representatives or and that no application for ign to the United States of the polytomia of the country of the country of the total country of the country of th
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Insert Provisional	(Number) I hereby claim the benefication(s) listed below.		States Code, §119(e) of any	
Application(s):	(Application Number)		(Filing	Date)
7,	(A. L. C. N		(Filing	(Date)
Insert Requested Information:	(Application Number) All Foreign Applications, if Months for Designs) Prior	To The Filing Date of Th	eventor's Certificate Filed his Application: pplication No.	More Than 12 Months (6 Date of Filing (Month/Day/Year)
(if appropriate) Insert Prior U.S.	I hereby claim the benefitsted below and, insofar as the prior United States application §112, I acknowledge the duty Code of Federal Regulations and the national or PCT interests.	ne subject matter of each on in the manner provided to disclose information v., §1.56 which became avarrational filing date of the	I by the first paragraph of I which is material to patental ailable between the filing dates application:	ition is not disclosed in the itle 35, United States Code, bility as defined in Title 37, ate of the prior application
Application(s):	(Application Number)	(Filing Date)	(Status - pa	tented, pending, abandoned)

(Filing Date)

(Application Number)

I hereby appoint the following attorneys to prosecute this application and/or an international application based on this application and to transact all business in the Patent and Trademark Office connected therewith and in connection with the resulting patent based on instructions received from the entity who first sent the application papers to the attorneys identified below, unless the inventor(s) or assignee provides said attorneys with a written notice to the contrary:

Terrell C. Birch	(Reg. No. 19,382)	Raymond C. Stewart	(Reg. No. 21,066)
Joseph A. Kolasch	(Reg. No. 22,463)	James M. Slattery	(Reg. No. <u>28,380</u>)
Bernard L. Sweeney	(Reg. No. 24,448)	Michael K. Mutter	(Reg. No. 29,680)
Charles Gorenstein	(Reg. No. 29,271)	Gerald M. Murphy, Jr.	(Reg. No. 28,977)
Leonard R. Svensson	(Reg. No. 30,330)	Terry L. Clark	(Reg. No. 32,644)
Andrew D. Meikle	(Reg. No. 32,868)	Marc S. Weiner	(Reg. No. 32,181)
Joe McKinney Muncy	(Reg. No. 32,334)	Andrew F. Reish	(Reg. No. <u>33,443</u>)
C. Joseph Faraci	(Reg. No. 32,350)	Donald J. Daley	(Reg. No. 34,313)
<i>J</i> ,	(C.

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Full Name of Second

Full Name of Third Inventor, if any

Full Name of Fourth

Full Name of Fifth Inventor, if any

Inventor, if any:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

			(month/day/year)
GIVEN NAME FAMILY NAME Keiichi TANAKA	INVENTOR'S SIGNATURE Keriche Tane		DATE* . 8/30/2001
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POST OFFICE ADDRESS (Complete Street Address c/o Oita University of 700	Dannoharu, Oita-sh	i, Oita, Ja	npan
GIVEN NAME Muhammad Shariq Vohra	INVENTOR'S SIGNATURE	,	DATE*
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GIVEN NAME FAMILY NAME	INVENTOR'S SIGNATURE		DATE*
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POST OFFICE ADDRESS (Complete Street Address	s including City, State & Country)		
GIVEN NAME FAMILY NAME	INVENTOR'S SIGNATURE		DATE*
Residence (City, State & Country)		CITIZENSHIP	
POST OFFICE ADDRESS (Complete Street Address	s including City, State & Country)		
GIVEN NAME FAMILY NAME	INVENTOR'S SIGNATURE		DATE*
Residence (City, State & Country)		CITIZENSHIP	
POST OFFICE ADDRESS (Complete Street Address	s including City, State & Country)	1	

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DATE OF SIGNATURE

BIRCH, STEWART, KOLASCH & BIRCH, LLP

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COMBINED DECLARATION AND POWER OF ATTORNEY

ATTORNEY DOCKET NO. 234-421P

YOU MUST
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FOR PATENT AND DESIGN APPLICATIONS As a below named inventor, I hereby declare that: my residence, post office address and citizenship are as stated next to my name; that I verily believe that I am the original, first and sole inventor (if only one inventor

Insert Title:	is named below) or an original matter which is claimed and for PRODUCTION OF	r which a patent is soi	ight on the invention ϵ	entitled:	:low) of the	subject
Fill in Appropriate Information -	the specification of which is att	ached hereto. If not	attached hereto,	,	26	
For Use Without Specification		was filed on ation Number		; and	_ as /or	
Attached:	the specification	was filed on	March 29, 2000		_as PCT	
	International Applic	ation Number	PCT/JP00/01959	; and	l was	
	amended under PCI	Article 19 on		(if applica	able)	
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Information:	Prior Foreign Application	(s)			Priority (Claimed
(if appropriate)	11-353257	Japan Japan	12/13/19 (Month/Day/Year		∑ Vaa	□ No
Ã.	(Number)	(Country)	(Month) Day, Tear	Theu,	Yes	No
•	(Number)	(Country)	(Month/Day/Year	Filed)	Yes	No
		(Country)	(Month/Day/Year	· Filed)	☐ Yes	□ No
•	(Number)	(Country)	(Month) Day, Tear	Tikeu)		
	(Number)	(Country)	(Month/Day/Year	Filed)	Yes	No
		(0	(Month/Day/Year	r Filed)	□ Yes	□ No
	(Number) I hereby claim the benefi	(Country)				
	application(s) listed below.	t under Tide 35, Offi	eu states coue, g115(c) of any onne	sa sattos pr	0110101111
Insert Provisional Application(s):	application (c) notes of the					
(if any)	(Application Number)			(Filing Date)		
	(Application Number)			(Filing Date)		
	All Foreign Applications, if a	env. for any Patent o	r Inventor's Certificat	e Filed More	Than 12 M	onths (6
Insert Requested Information: (if appropriate)	Months for Designs) Prior T	o The Filing Date of	This Application: Application No.		f Filing (Month	
	I hereby claim the benefi listed below and, insofar as th prior United States applicatio §112, I acknowledge the duty Code of Federal Regulations, and the national or PCT inter	te subject matter of ea in the manner provi to disclose information §1.56 which became	ch of the claims of this ded by the first paragra on which is material to j available between the	s application is aph of Title 35, patentability a	s not disclos , United Star s defined in	ed in the tes Code, Title 37,

(Filing Date)

(Filing Date)

Page 1 of 2

(if any)

Insert Prior U.S. Application(s):

(Application Number)

(Application Number)

(Status - patented, pending, abandoned)

(Status - patented, pending, abandoned)

I hereby appoint the following attorneys to prosecute this application and/or an international application based on this application and to transact all business in the Patent and Trademark Office connected therewith and in connection with the resulting patent based on instructions received from the entity who first sent the application papers to the attorneys identified below, unless the inventor(s) or assignee provides said attorneys with a written notice to the contrary:

	,	4	
Terrell C. Birch	(Reg. No. 19,382)	Raymond C. Stewart	(Reg. No. 21,066)
Joseph A. Kolasch	(Reg. No. 22,463)	James M. Slattery	(Reg. No. 28,380)
Bernard L. Sweeney	(Reg. No. 24,448)	Michael K. Mutter	(Reg. No. 29,680)
Charles Gorenstein	(Reg. No. 29,271)	Gerald M. Murphy, Jr.	(Reg. No. 28,977)
Leonard R. Svensson	(Reg. No. 30,330)	Terry L. Clark	(Reg. No. 32,644)
Andrew D. Meikle	(Reg. No. 32,868)	Marc S. Weiner	(Reg. No. 32,181)
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Full Name of Fourth Inventor, if any

Full Name of Fifth Inventor, if any

see above

see above

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

				(month/day/year)				
GIVEN NAME Keiichi T		INVENTOR'S SIGNATURE		DATE*				
· · · · · · · · · · · · · · · · · · ·	Oita, Japan		CITIZENSHIP	Japan				
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GIVEN NAME Muhammad	FAMILY NAME Shariq Vohra	INVENTOR'S SIGNATURE Mahammad Shaxqishx	<i>(-2,</i>	DATE* 9/5/2001				
	Ontario, Canada		CITIZENSHIP	Pakistan				
2	, ,	es including City, State & Country) Dean, Ontario, Cana	da					
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE		DATE*				
Residence (City, Sta	te & Country)		CITIZENSHIP	<u> </u>				
POST OFFICE ADDI	RESS (Complete Street Addre	ss including City, State & Country)						
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE		DATE*				
Residence (City, Sta	te & Country)		CITIZENSHIP					
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Residence (City, Sta	ite & Country)		CITIZENSHIP					
POST OFFICE ADDI	RESS (Complete Street Addre	ess including City, State & Country)	<u> </u>					
* DATE OF SIGNATU	RF .		*					

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